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					Date of Report PRISCA	23-01-20 5.0.2.37
Patient Data						
Name	М	RS SEEN	MA SHARMA	Patient ID		062001220004
Birthday			30-12-90) Sample ID		10511154
Age at delivery			29.	Sample Date		22/01/2020
Gestational age			13+0	5		
Correction factors						
Fetuses	1 1	VF		unknown	Previous trisomy 21	unknown
Weight in kg	73]	Diabetes		unknown	Pregnancies	unknown
Smoker	Unknown	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational ag	5	13+5
PAPP-A	3.85 1	nIU/ml	0.69	Method		CRL(<>Robinson
fb-hCG	37.5 1	ıg/ml	0.99	Scan date		21-01-20
Risks at sampling date				Crown rump length in mm 65		
Age Risk			1:752	Nuchal translu	cency MOM	0.9
Biochemical T21 risk			1:2058	Nasal bone		Present
Combined Trisomy 21	Risk		<1:10000	Sonographer		DR.SAGAR
Trisomy 13/18 + NT			<1:10000	Qualification i	n measuring NT	MBBS,MD
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
1:10 1:100 1:250 Cut off 1:1000 1:1000			The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 women with not affected pregnancies. The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that the risk calculations are statistical aapproaches and have no diagnostic value!			
13 15 17 19 21 23 25 Trisomy 13/18 + NT The calculated risk for <1:10000, which indic	r Trisomy 13	/18 (with k	Age	The laborator	y cannot be hold resp essment! Calculated 1	oonsible for their impact risks have no diagnostic Risk below Age risk